

Refine Search

Search Results -

Term	Documents
(9 NOT 10).PGPB,USPT,USOC,EPAB,JPAB,DWPI,TDBD.	44
(L9 NOT L10 ).PGPB,USPT,USOC,EPAB,JPAB,DWPI,TDBD.	44

Database:

US Pre-Grant Publication Full-Text Database  
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IBM Technical Disclosure Bulletins

Search:

L11

Refine Search

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DATE: Wednesday, April 05, 2006    [Printable Copy](#)    [Create Case](#)

<u>Set</u> <u>Name</u> side by side	<u>Query</u>	<u>Hit</u> <u>Count</u>	<u>Set</u> <u>Name</u> result set
DB=PGPB,USPT,USOC,EPAB,JPAB,DWPI,TDBD; THES=ASSIGNEE; PLUR=YES; OP=AND			
<u>L11</u>	L9 not L10	44	<u>L11</u>
<u>L10</u>	L9 and (PDGF or PDGF-AA or PDGF-AB or PDGF-BB)	8	<u>L10</u>
<u>L9</u>	L8 not L5	52	<u>L9</u>
<u>L8</u>	L7 and L6	56	<u>L8</u>
<u>L7</u>	(Promoter) same (lactalbumin or casein or lactoglobulin or (mammary adj epithelial))	1552	<u>L7</u>
<u>L6</u>	(transgenic adj animal) same (bioreactor)	297	<u>L6</u>
<u>L5</u>	L3 and (transgenic adj animal)	32	<u>L5</u>
<u>L4</u>	L3 and (transgenic adj (mammal or mouse or rat or bovine or ovine or porcine or caprine or equine or buffalo))	0	<u>L4</u>
<u>L3</u>	(PDGF or PDGF-AA or PDGF-AB or PDGF-BB) same (milk)	90	<u>L3</u>

L2 L1 and PDGF  
L1 Echelard-Yann.in.

1 L2  
20 L1

END OF SEARCH HISTORY

## Welcome to DialogClassic Web(tm)

Dialog level 05.10.03D  
Last logoff: 04apr06 15:40:16  
Logon file001 05apr06 09:44:06

## \*\*\* ANNOUNCEMENTS \*\*\*

\*\*\*

## NEW FILES RELEASED

\*\*\*Regulatory Affairs Journals (File 183)  
\*\*\*Index Chemicus (File 302)  
\*\*\*Inspec (File 202)

\*\*\*

## RELOADS COMPLETED

\*\*\* MEDLINE has been reloaded with the 2006 MeSH (Files 154 & 155)  
\*\*\* The 2005 reload of the CLAIMS files (Files 340, 341, 942)  
is now available online.

## RESUMED UPDATING

\*\*\*EDGARPLUS(TM)-Williams Act Filings (File 773)  
\*\*\*EDGARPLUS(TM)-Prospectuses (File 774)  
\*\*\*EDGARPLUS(TM)-Registration Statements (File 775)  
\*\*\*EDGARPLUS(TM)-6K,8K, and 10C Filings (File 776)  
\*\*\*EDGARPLUS(TM)-10-K & 20F Filings (File 778)  
\*\*\*EDGARPLUS(TM)-10-Q Filings (File 779)  
\*\*\*EDGARPLUS(TM)-Proxy Statements (File 780)

\*\*\*

Chemical Structure Searching now available in Prous Science Drug Data Report (F452),  
IMS R&D Focus (F445/955), Pharmaprojects (F128/928), Beilstein  
Facts (F390), Derwent Chemistry Resource (F355) and Index Chemicus  
(File 302).

\*\*\*

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>>><http://www.dialog.com/whatsnew/>. You can find news about<<<  
>>>a specific database by entering HELP NEWS <file number>.<<<  
KWIC is set to 50.

HIGHLIGHT set on as ' '

\* \* \*

File 1:ERIC 1966-2006/Feb  
(c) format only 2006 Dialog

Set Items Description

--- -----

Cost is in DialUnits  
?

B 155, 5, 73

05apr06 09:44:18 User259876 Session D860.1

\$0.81 0.230 DialUnits File1

\$0.81 Estimated cost File1

\$0.05 INTERNET

\$0.86 Estimated cost this search

\$0.86 Estimated total session cost 0.230 DialUnits

SYSTEM:OS - DIALOG OneSearch

File 155:MEDLINE(R) 1951-2006/Apr 05

(c) format only 2006 Dialog

\*File 155: Medline has been reloaded. Some accession numbers  
have changed.

File 5:Biosis Previews(R) 1969-2006/Apr W1

(c) 2006 BIOSIS  
 File 73:EMBASE 1974-2006/Apr 04  
 (c) 2006 Elsevier Science B.V.

Set	Items	Description
---	-----	-----
?		
S	(PDGF OR PDGF-AA OR PDGF-AB OR PDGF-BB) (S) (MILK OR TRANSGENIC OR BIOREACTOR)	
	25922	PDGF
	16	PDGF-AA
	16	PDGF-AB
	90	PDGF-BB
	214055	MILK
	170029	TRANSGENIC
	22846	BIOREACTOR
S1	292	(PDGF OR PDGF-AA OR PDGF-AB OR PDGF-BB) (S) (MILK OR TRANSGENIC OR BIOREACTOR)
?		
S	(PROMOTER) (S) (LACTALBUMIN OR CASEIN OR LACTOGLOBULIN OR (MAMMARY (W) EPITHELIAL)	
	344531	PROMOTER
	8471	LACTALBUMIN
	57760	CASEIN
	8200	LACTOGLOBULIN
	145691	MAMMARY
	465062	EPITHELIAL
	11966	MAMMARY(W)EPITHELIAL
S2	2123	(PROMOTER) (S) (LACTALBUMIN OR CASEIN OR LACTOGLOBULIN OR (MAMMARY (W) EPITHELIAL))
?		
S S1 AND S2		
	292	S1
	2123	S2
S3	0	S1 AND S2
?		
Set	Items	Description
S1	292	(PDGF OR PDGF-AA OR PDGF-AB OR PDGF-BB) (S) (MILK OR TRANSGENIC OR BIOREACTOR)
S2	2123	(PROMOTER) (S) (LACTALBUMIN OR CASEIN OR LACTOGLOBULIN OR (MAMMARY (W) EPITHELIAL))
S3	0	S1 AND S2
?		
S S1 AND (TRANSGENIC (W) (ANIMAL OR MOUSE OR RAT OR BOVINE OR GOAT OR PIG))		
	292	S1
	170029	TRANSGENIC
	3384884	ANIMAL
	1739122	MOUSE
	2944679	RAT
	431332	BOVINE
	51861	GOAT
	426399	PIG
	44763	TRANSGENIC(W) ((( (ANIMAL OR MOUSE) OR RAT) OR BOVINE) OR GOAT) OR PIG)
S4	82	S1 AND (TRANSGENIC (W) (ANIMAL OR MOUSE OR RAT OR BOVINE OR GOAT OR PIG))

?

S S4 NOT PY&gt;2000

82 S4

8397784 PY&gt;2000

S5 31 S4 NOT PY&gt;2000

?

RD

S6 27 RD (unique items)

?

Set	Items	Description
S1	292	(PDGF OR PDGF-AA OR PDGF-AB OR PDGF-BB) (S) (MILK OR TRANS-GENIC OR BIOREACTOR)
S2	2123	(PROMOTER) (S) (LACTALBUMIN OR CASEIN OR LACTOGLOBULIN OR - (MAMMARY (W) EPITHELIAL))
S3	0	S1 AND S2
S4	82	S1 AND (TRANSGENIC (W) (ANIMAL OR MOUSE OR RAT OR BOVINE OR GOAT OR PIG))
S5	31	S4 NOT PY>2000
S6	27	RD (unique items)

?

T S6/3,K/ALL

6/3,K/1 (Item 1 from file: 155)

DIALOG(R) File 155:MEDLINE(R)

(c) format only 2006 Dialog. All rts. reserv.

11556032 PMID: 9396344

**[Molecular pathomechanism of HTLV-I infectious diseases]**

Kitajima I

Department of Laboratory Medicine, Kagoshima University of School of Medicine.

Rinsho byori. The Japanese journal of clinical pathology (JAPAN) Nov 1997, 45 (11) p1048-56, ISSN 0047-1860--Print Journal Code: 2984781R

Publishing Model Print

Document type: Journal Article; Review ; English Abstract

Languages: JAPANESE

Main Citation Owner: NLM

Record type: MEDLINE; Completed

...detected mRNA for HTLV-I tax/rex in cultured synovial cells by reverse transcription polymerase chain reaction. Moreover, induction of chronic inflammatory arthropathy in mice **transgenic** for HTLV-I tax gene strongly suggested the pathogenic mechanism of HAAP. Histologic findings of affected joints in mice showed erosions of bones and pannus-like granulomatous change with infiltration of mononuclear cells. Thus, this novel mechanism might explain synovial proliferation caused by HTLV-I. Tax-expressing **transgenic mouse** lines also demonstrated that tax itself could serve as an oncogene in fibroblastic cells. Tumors occurred in 100% of the mice with reproducible time periods after wounding. We established cell lines, which expressed high levels of c-fos, c-myc, myb, **PDGF**, TGF-beta, Zif, and IL-6. Antisense ablation of the p65 subunits of NF-kappa B profoundly inhibited tumor growth in vitro with no apparent affect on the growth of normal cells. These studies were successfully extended to tax- **transgenic** animals. Intraperitoneal injections of NF-kappa B p65 antisense at the 40 micrograms/g weight dose led to growth arrest after 7 days, and apparent...

6/3,K/2 (Item 2 from file: 155)

DIALOG(R)File 155:MEDLINE(R)

(c) format only 2006 Dialog. All rts. reserv.

11535030 PMID: 9368101

**Temporal and spatial specificity of PDGF alpha receptor promoter in transgenic mice.**

Reinertsen K K; Bronson R T; Stiles C D; Wang C

Department of Microbiology and Molecular Genetics, Harvard Medical School, Boston, MA, USA.

Gene expression (UNITED STATES) 1997, 6 (5) p301-14, ISSN 1052-2166--Print Journal Code: 9200651

Contract/Grant No.: CA 74907; CA; NCI; HD 24926; HD; NICHD

Publishing Model Print; Erratum in Gene Expr 1998;7(2) 131

Document type: Journal Article

Languages: ENGLISH

Main Citation Owner: NLM

Record type: MEDLINE; Completed

**Temporal and spatial specificity of PDGF alpha receptor promoter in transgenic mice.**

Aberrant expression of the platelet-derived growth factor alpha receptor (PDGF alpha R) has been linked to developmental abnormalities in vertebrate models, and has been implicated in multiple disease states in humans. To identify cis-acting regulatory elements that dictate expression of this receptor, we generated **transgenic** mice bearing the reporter gene beta-galactosidase (lacZ) under the control of a 6-kb promoter sequence. Expression of lacZ was monitored throughout embryonic development, with special focus on nervous tissue, skeleton, and several organ systems wherein PDGF alpha R expression is thought to play a pivotal role. In several independent **transgenic mouse** strains, lacZ expression recapitulated predominant features of PDGF alpha R gene expression during mouse development. These results demonstrate that critical tissue-specific regulatory elements for PDGF alpha R expression are located within a 6-kb upstream region of the PDGF alpha R gene.

6/3,K/3 (Item 3 from file: 155)

DIALOG(R)File 155:MEDLINE(R)

(c) format only 2006 Dialog. All rts. reserv.

08961113 PMID: 1916630

**[HTLV-I tax mediated activation of cellular genes in transgenic mice]**

Shinohara T

Second Department of Pathology, Hokkaido University School of Medicine, Sapporo, Japan.

Hokkaido igaku zasshi The Hokkaido journal of medical science (JAPAN) Jul 1991, 66 (4) p534-43, ISSN 0367-6102--Print Journal Code: 17410290R

Publishing Model Print

Document type: Journal Article ; English Abstract

Languages: JAPANESE

Main Citation Owner: NLM

Record type: MEDLINE; Completed

... tax have been studied in vitro, mostly in T-cell lines. To determine its function in vivo in multiple cell types, we have used two **transgenic mouse** lines in which tax is expressed under the control of the LTR

(LTRtax) or murine Thyl. 2 (Thytax) transcriptional regulatory sequences. Tax protein is expressed...

... gland, skeletal muscle, bone matrix and thymus tissue. In these tissues the expression of endogenous IL-2R, c-fos, GM-CSF, Zif268, IL-6, and PDGF -B were studied. In fibroblastic tumors GM-CSF, IL-6, PDGF -B, Zif268, c-fos were expressed at high levels. No significant changes in expression of these genes were seen in other tissues. This suggests that...

6/3,K/4 (Item 1 from file: 5)  
DIALOG(R)File 5:Biosis Previews(R)  
(c) 2006 BIOSIS. All rts. reserv.

0012937450 BIOSIS NO.: 200100109289

**Polyglobulia in transgenic mice overexpressing erythropoietin worsens outcome after focal brain ischemia**

AUTHOR: Wiessner C (Reprint); Allegrini P R; Alt U R; Ekatodramis D; Gassmann M

AUTHOR ADDRESS: Novartis Pharma AG, Basel, Switzerland\*\*Switzerland

JOURNAL: Society for Neuroscience Abstracts 26 (1-2): pAbstract No.-670.11  
2000 2000

MEDIUM: print

CONFERENCE/MEETING: 30th Annual Meeting of the Society of Neuroscience New Orleans, LA, USA November 04-09, 2000; 20001104

SPONSOR: Society for Neuroscience

ISSN: 0190-5295

DOCUMENT TYPE: Meeting; Meeting Abstract

RECORD TYPE: Abstract

LANGUAGE: English

ABSTRACT: **Transgenic mouse** lines expressing human Erythropoietin (EPO) under the control of the **PDGF** promoter were investigated in a stroke model (permanent MCAO). In line tg6, CNS and serum EPO levels were increased, resulting in a hematocrit about 80...

6/3,K/5 (Item 2 from file: 5)  
DIALOG(R)File 5:Biosis Previews(R)  
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0012910059 BIOSIS NO.: 200100081898

**Cytoplasmatic and nuclear localization of ataxin-7 (a7) in normal human brain and nuclear accumulation of mutant a7 in transgenic mouse models of SCA 7**

AUTHOR: Lindenberg K S (Reprint); Devys D; Mueller K; Landwehrmeyer G B; Mandel J L; Volk B; Weber C; Yvert G

AUTHOR ADDRESS: U. Freiburg, Freiburg, Germany\*\*Germany

JOURNAL: Society for Neuroscience Abstracts 26 (1-2): pAbstract No.-479.7  
2000 2000

MEDIUM: print

CONFERENCE/MEETING: 30th Annual Meeting of the Society of Neuroscience New Orleans, LA, USA November 04-09, 2000; 20001104

SPONSOR: Society for Neuroscience

ISSN: 0190-5295

DOCUMENT TYPE: Meeting; Meeting Abstract

RECORD TYPE: Abstract

LANGUAGE: English

**Cytoplasmatic and nuclear localization of ataxin-7 (a7) in normal human**

**brain and nuclear accumulation of mutant a7 in transgenic mouse models of SCA 7**

...ABSTRACT: these neurons are vulnerable in SCA7, a physiological nuclear enrichment of a7 may predispose to neurodegeneration. To gain further insight into the pathogenesis of SCA7, **transgenic mouse** models were generated by using **PDGF** -B or pcp-2 as promoters to drive the expression of full length mutant or normal a7 in neurons throughout the brain or in Purkinje...

**6/3,K/6 (Item 3 from file: 5)**

DIALOG(R)File 5:BIOSIS Previews(R)  
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0012240577 BIOSIS NO.: 199900500237

**Platelet derived growth factor-AA ( PDGF -AA) in liver fibrosis: An inducible transgenic mouse model to study liver fibrogenesis**

AUTHOR: Kanzler Stephan (Reprint); Blessing Manfred; Galle Peter R; Lohse Ansgar W

AUTHOR ADDRESS: University of Mainz, Mainz, Germany\*\*Germany

JOURNAL: Hepatology 30 (4 PART 2): p413A Oct., 1999 1999

MEDIUM: print

CONFERENCE/MEETING: 50th Annual Meeting and Postgraduate Courses of the American Association for the Study of Liver Diseases Dallas, Texas, USA November 5-9, 1999; 19991105

SPONSOR: American Association for the Study of Liver Diseases

ISSN: 0270-9139

DOCUMENT TYPE: Meeting; Meeting Abstract

RECORD TYPE: Citation

LANGUAGE: English

**Platelet derived growth factor-AA ( PDGF -AA) in liver fibrosis: An inducible transgenic mouse model to study liver fibrogenesis**

**6/3,K/7 (Item 4 from file: 5)**

DIALOG(R)File 5:BIOSIS Previews(R)  
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0011513178 BIOSIS NO.: 199800307425

**Oligodendrocyte population dynamics and the role of PDGF in vivo**

AUTHOR: Calver Andrew R; Hall Anita C; Yu Wei-Ping; Walsh Frank S; Heath John K; Betsholtz Christer; Richardson William D (Reprint)

AUTHOR ADDRESS: MRC Lab. Molecular Cell Biol., Dep. Biol., Univ. Coll. London, Gower St., London WC1E 6BT, UK\*\*UK

JOURNAL: Neuron 20 (5): p869-882 May, 1998 1998

MEDIUM: print

ISSN: 0896-6273

DOCUMENT TYPE: Article

RECORD TYPE: Abstract

LANGUAGE: English

...ABSTRACT: rise to oligodendrocytes. Progenitor cell proliferation stops before birth because the cell cycle slows down, linked to an increase in differentiation and death. Experiments with **transgenic** mice show that platelet-derived growth factor ( **PDGF** ) drives progenitor cell division and suggest that slowing of and exit from the cycle reflects a decline in **PDGF** signaling. Overexpressing **PDGF** induces hyperproliferation of progenitor cells and excessive, ectopic production of oligodendrocytes.

However, the superfluous oligodendrocytes die at an immature stage of differentiation, leaving a normal...

## DESCRIPTORS:

ORGANISMS: **transgenic mouse** (Muridae)

6/3,K/8 (Item 5 from file: 5)

DIALOG(R)File 5:Biosis Previews(R)

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0011147584 BIOSIS NO.: 199799781644

**Urokinase and tissue-type plasminogen activator are required for the mitogenic and chemotactic effects of bovine fibroblast growth factor and platelet-derived growth factor-BB for vascular smooth muscle cells**

AUTHOR: Herbert Jean-Marc (Reprint); Lamarche Isabelle; Carmeliet Peter

AUTHOR ADDRESS: Haemobiology Res. Dep., Sanofi Recherche, 195 Route d'Espagne, 31036 Toulouse, France\*\*France

JOURNAL: Journal of Biological Chemistry 272 (38): p23585-23591 1997 1997

ISSN: 0021-9258

DOCUMENT TYPE: Article

RECORD TYPE: Abstract

LANGUAGE: English

...ABSTRACT: PA) and urokinase-type plasminogen activator (u-PA) in the mitogenic and chemotactic potential of bovine fibroblast growth factor (bFGF) and platelet-derived growth factor ( **PDGF** )-BB for smooth muscle cells (SMC). Aortic SMC were isolated from **transgenic** mice showing single inactivations of the t-PA, u-PA, plasminogen activator inhibitor-1, or urokinase-type plasminogen activator receptor (u-PAR) genes. With regard...

...serum-induced proliferation, all cell types showed similar responses. However, SMC isolated from t-PA-deficient mice did not proliferate or migrate in response to **PDGF** , whereas SMC isolated from u-PA-deficient animals appeared to be much less sensitive to bFGF than the cells isolated from the other animals. Supplementation...

...or in wild-type SMC, cultured in the presence of antibodies to u-PAR. The role of u-PA and t-PA in bFGF and **PDGF** -induced growth and migration of SMC was not dependent on plasmin generation and activity as demonstrated by the inactivity of epsilon-aminocaproic acid and aprotinin ...

...state levels of u-PA and t-PA mRNA and proteins were observed after 24 h of incubation of the cell cultures with bFGF and **PDGF** -BB, respectively. These results therefore indicate that, at least in vitro, t-PA is an important element of the activity of **PDGF** -BB with regard to the proliferation and migration of SMC whereas u-PA is a key factor in the effect of bFGF on SMC.

## DESCRIPTORS:

MISCELLANEOUS TERMS: ... **TRANSGENIC ANIMAL** ;

6/3,K/9 (Item 6 from file: 5)

DIALOG(R)File 5:Biosis Previews(R)

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0010856867 BIOSIS NO.: 199799490927

**Molecular and anatomic analysis of the PDGF -hAPP V717F transgenic mouse**

AUTHOR: Hyman Bradley T (Reprint); Irizarry Michael C; McNamara Megan;

Soriano Ferdi; Schenk Dale; Games Dora  
AUTHOR ADDRESS: Charlestown, MA, USA\*\*USA  
JOURNAL: Neurology 48 (3 SUPPL. 2): pA273 1997 1997  
CONFERENCE/MEETING: 49th Annual Meeting of the American Academy of  
Neurology Boston, Massachusetts, USA April 12-19, 1997; 19970412  
ISSN: 0028-3878  
DOCUMENT TYPE: Meeting; Meeting Abstract  
RECORD TYPE: Citation  
LANGUAGE: English

**Molecular and anatomic analysis of the PDGF -hAPP V717F transgenic  
mouse**

## DESCRIPTORS:

MISCELLANEOUS TERMS: ... PDGF -HAPP V717F TRANSGENIC ANIMAL MODEL

6/3,K/10 (Item 7 from file: 5)  
DIALOG(R)File 5:Biosis Previews(R)  
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0010581286 BIOSIS NO.: 199699215346

**Molecular and anatomic correlates in the PDGF -hAPP V717F transgenic  
mouse**

AUTHOR: Irizarry M C (Reprint); Page K J; Soriano F; Schnek D; Games D;  
Hyman B T  
AUTHOR ADDRESS: Neurol., Mass Gen. Hosp., Boston, MA 02114, USA\*\*USA  
JOURNAL: Society for Neuroscience Abstracts 22 (1-3): p25 1996 1996  
CONFERENCE/MEETING: 26th Annual Meeting of the Society for Neuroscience  
Washington, D.C., USA November 16-21, 1996; 19961116  
ISSN: 0190-5295  
DOCUMENT TYPE: Meeting; Meeting Abstract; Meeting Slide  
RECORD TYPE: Citation  
LANGUAGE: English

**Molecular and anatomic correlates in the PDGF -hAPP V717F transgenic  
mouse**

6/3,K/11 (Item 8 from file: 5)  
DIALOG(R)File 5:Biosis Previews(R)  
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0010069178 BIOSIS NO.: 199598537011

**Role of platelet-derived growth factor ( PDGF ) in hepatic fibrosis:  
Evaluation in a novel transgenic mouse model**

AUTHOR: Davern T J (Reprint); Liao X; Ferrell L; Rockey D (Reprint);  
Friedman S L (Reprint); Escabedo J A; Williams L T; Scharschmidt B F  
(Reprint)  
AUTHOR ADDRESS: UCSF Liver Cent., Univ. Calif., San Francisco, CA 94143,  
USA\*\*USA  
JOURNAL: Hepatology 22 (4 PART 2): p281A 1995 1995  
CONFERENCE/MEETING: 46th Annual Meeting and Postgraduate Course of the  
American Association for the Study of Liver Diseases Chicago, Illinois,  
USA November 3-7, 1995; 19951103  
ISSN: 0270-9139  
DOCUMENT TYPE: Meeting; Meeting Abstract  
RECORD TYPE: Citation  
LANGUAGE: English

**Role of platelet-derived growth factor ( PDGF ) in hepatic fibrosis:**

**Evaluation in a novel transgenic mouse model**

6/3,K/12 (Item 1 from file: 73)

DIALOG(R)File 73:EMBASE

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11526704 EMBASE No: 2002098495

**Experimental models of growth factor-mediated angiogenesis and blood-retinal barrier breakdown**

Vinores S.A.; Seo M.S.; Okamoto N.; Ash J.D.; Wawrousek E.F.; Xiao W.-H.; Hudish T.; Derevjani N.L.; Campochiaro P.A.

S.A. Vinores, Wilmer Eye Institute, Johns Hopkins Univ. School of Med.,  
825 Maumenee Building, 600 North Wolfe Street, Baltimore, MD 21287-9289  
United States

AUTHOR EMAIL: svinores@jhmi.edu

General Pharmacology: Vascular System ( GEN. PHARMACOL. VASC. SYST. ) ( United States) 2000, 35/5 (233-239)

CODEN: GEPHD ISSN: 0306-3623

PUBLISHER ITEM IDENTIFIER: S0306362301001173

DOCUMENT TYPE: Journal ; Conference Paper

LANGUAGE: ENGLISH SUMMARY LANGUAGE: ENGLISH

NUMBER OF REFERENCES: 73

...results in neovascularization (NV) that originates from the vascular bed closest to the ganglion cell layer. To study the effects of VEGF, independent lines of **transgenic** mice that express VEGF in the lens and in the retina have been generated. Expression in the lens results in excessive proliferation and accumulation of...

...blood vessel organization or maturation in the prenatal mouse. Abnormal vessels do form on the retinal surface, but not until the second postnatal week. In **transgenic** mice expressing VEGF in the photoreceptors, NV originates from the deep capillary bed - the vascular bed closest to the photoreceptors. NV is accompanied by localized blood-retinal barrier breakdown. NV is also induced in **PDGF** -B **transgenic** mice.  $\square$ PDGF $\square$  -B expression in the lens occurs prenatally and, during this time, mainly affects the perilenticular vessels. Postnatally, **transgenic** mice expressing **PDGF** -B in the lens or photoreceptors show a similar phenotype. In both models, a highly vascularized cell mass containing endothelial cells, pericytes, and glia forms...

**MEDICAL DESCRIPTORS:**

**transgenic mouse** ; protein expression; lens; retina; cell proliferation; endothelium cell; blood vessel; prenatal period; surface property; photoreceptor; phenotype; cell assay; pericyte; glia; vascularization; retina blood vessel; cell...

6/3,K/13 (Item 2 from file: 73)

DIALOG(R)File 73:EMBASE

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11051916 EMBASE No: 2000398852

**Articular cartilage and growth factors (first part)**

CARTILAGO ARTICULAR Y FACTORES DE CRECIMIENTO (PRIMERA PARTE)

Vega J.A.; Garcia-Suarez O.; Martinez-Almagro A.

J.A. Vega, Depto. de Morfol. y Biologia Celular, Facultad de Medicina, C/  
Julian Claveria, s/n, 33006 Oviedo Spain

Mapfre Medicina ( MAPFRE MED. ) (Spain) 2000, 11/3 (212-225)

CODEN: MAMEE ISSN: 1130-5665

DOCUMENT TYPE: Journal ; Review  
 LANGUAGE: SPANISH SUMMARY LANGUAGE: ENGLISH; SPANISH  
 NUMBER OF REFERENCES: 120

...have been used successfully in the experimental treatment of some of them. This paper is a review of the role of IGFs, TFGs, FGFs, EGF, **PDGF** and neurotrophins growth factors and cytokines. Data obtained from **transgenic** animals and the effects genetic therapy using transfected chondrocytes as vectors for the genes involved in the cartilage biology are also considered.

MEDICAL DESCRIPTORS:

cartilage cell; cell survival; extracellular matrix; gene therapy;  
**transgenic animal** ; human; nonhuman; human tissue; review

6/3,K/14 (Item 3 from file: 73)  
 DIALOG(R)File 73:EMBASE  
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11001897 EMBASE No: 2001045234

**Photoreceptor-specific expression of platelet-derived growth factor-B results in traction retinal detachment**

Man Seong Seo; Okamoto N.; Vinores M.A.; Vinores S.A.; Hackett S.F.; Yamada H.; Yamada E.; Derevjani N.L.; LaRochelle W.; Zack D.J.; Campochiaro P.A.

Dr. P.A. Campochiaro, Maumenee 719, Johns Hopkins Univ. Sch. of Medicine, 600 N. Wolfe Street, Baltimore, MD 21287-9277 United States

AUTHOR EMAIL: pcampo@jhmi.edu

American Journal of Pathology ( AM. J. PATHOL. ) (United States) 2000, 157/3 (995-1005)

CODEN: AJPAA ISSN: 0002-9440

DOCUMENT TYPE: Journal ; Article

LANGUAGE: ENGLISH SUMMARY LANGUAGE: ENGLISH

NUMBER OF REFERENCES: 42

Expression of platelet-derived growth factor ( **PDGF** )-A and **PDGF** -B is increased in patients with proliferative retinopathies in which traction retinal detachments occur. Previous studies have demonstrated that increased expression of **PDGF** -A in the retina of **transgenic** mice results in retinal gliosis due to proliferation of astrocytes with different retinal phenotypes based on the time of onset and location of the **PDGF** -A production. In this study, we investigated the effects of **PDGF** -B in the retina using gain-of-function **transgenic** mice that express **PDGF** -B in photoreceptors. These mice show proliferation of astrocytes, pericytes, and, to a lesser extent, endothelial cells, resulting in ectopic cells on the surface and...

...of cells exert traction on the retina resulting in traction retinal detachments similar to those seen in humans with proliferative retinopathies. These studies suggest that **PDGF** -B has more dramatic effects in the retina than **PDGF** -A, because it acts on additional cell types, in particular on pericytes, which have a highly developed contractile apparatus. These studies in the retina suggest a means that could be used in other tissues throughout the body to achieve graded **PDGF** effects. They also provide a new model of traction retinal detachment that can be used to investigate new treatments for patients with proliferative retinopathies.

MEDICAL DESCRIPTORS:

proliferative retinopathy--diagnosis--di; **transgenic mouse** ; gliosis; astrocyte; cell proliferation; phenotype; onset age; protein localization;

pericyte; endothelium cell; cell type; nonhuman; mouse; animal experiment;  
animal model; controlled study; animal tissue; animal...

6/3,K/15 (Item 4 from file: 73)

DIALOG(R)File 73:EMBASE

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10947989 EMBASE No: 2000439936

**The role of platelet-derived growth factor in a murine model of crescentic nephritis**

Haseley L.A.; Pippin J.W.; Huang X.R.; Lan H.Y.; Gordon K.L.; Seifert R.A.; Johnson R.J.

L.A. Haseley, Box 356521, Division of Nephrology, Univ. of Washington Medical Center, Seattle, WA 98195 United States

Nephrology ( NEPHROLOGY ) (Australia) 2000, 5/3 (193-199)

CODEN: NEPHF ISSN: 1320-5358

DOCUMENT TYPE: Journal; Article

LANGUAGE: ENGLISH SUMMARY LANGUAGE: ENGLISH

NUMBER OF REFERENCES: 38

Platelet-derived growth factor ( **PDGF** ) is a major mesenchymal cell mitogen, with an established role in the pathogenesis of experimental mesangial proliferative nephritis. The role of **PDGF** in experimental models of crescentic glomerulonephritis is not well defined. To study the role of **PDGF** in glomerular crescent formation, we induced a model of crescentic glomerulonephritis in **transgenic** mice expressing high concentrations of the soluble external domain of the PDGFbeta receptor ( **PDGF** -Rbeta). Crescentic nephritis was induced by the intraperitoneal injection of antibody to whole rabbit glomeruli. At day 7 of disease, biopsies of **transgenic** and wild-type mice were evaluated for crescent frequency, crescent area, and thickness of crescent cell layer. In situ hybridization was performed to evaluate the expression of both **PDGF** B-chain and PDGFRbeta mRNA within crescents. Delivery of soluble receptor to the urinary space was evaluated by Western blotting. Crescent frequency did not differ between wild type and **transgenic** mice. However, crescent area quantified by computer image analysis was significantly reduced in **transgenic** mice (P<0.015). **Transgenic** biopsies displayed predominantly crescents composed of two cell layers (P=0.03 compared with wild type), whereas wild-type biopsies had significantly more crescents composed of four or more cell layers (P=0.04). Both **PDGF** B-chain and **PDGF** -RbetamRNA were detected within crescents in a heterogeneous fashion. Soluble receptor was detectable in the urine of all **transgenic** diseased mice. We conclude that **PDGF** plays a role in modulating crescent size and development in our murine model of crescentic nephritis.

**MEDICAL DESCRIPTORS:**

pathogenesis; **transgenic** mouse ; gene expression; kidney biopsy;  
mesenchyme; glomerulus; nonhuman; mouse; animal experiment; animal model;  
controlled study; animal tissue; article; priority journal

6/3,K/16 (Item 5 from file: 73)

DIALOG(R)File 73:EMBASE

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10886103 EMBASE No: 2000370690

**PDGF-C is a new protease-activated ligand for the PDGF alpha-receptor**

Li X.; Ponten A.; Aase K.; Karlsson L.; Abramsson A.; Uutela M.; Backstrom G.; Hellstrom M.; Bostrom H.; Li H.; Soriano P.; Betsholtz C.; Heldin C.-H.; Alitalo K.; Ostman A.; Eriksson U.

U. Eriksson, Ludwig Institute for Cancer Research, Stockholm Branch, Box 240, S-17177 Stockholm Sweden  
 AUTHOR EMAIL: ueri@licr.ki.se  
 Nature Cell Biology ( NATURE CELL BIOL. ) (United Kingdom) 2000, 2/5 (302-307)  
 CODEN: NCBIF ISSN: 1465-7392  
 DOCUMENT TYPE: Journal; Article  
 LANGUAGE: ENGLISH SUMMARY LANGUAGE: ENGLISH  
 NUMBER OF REFERENCES: 43

Platelet-derived growth factors (PDGFs) are important in many types of mesenchymal cell. Here we identify a new **PDGF** , **PDGF** -C, which binds to and activates the **PDGF** alpha-receptor. **PDGF** -C is activated by proteolysis and induces proliferation of fibroblasts when overexpressed in **transgenic** mice. In situ hybridization analysis in the murine embryonic kidney shows preferential expression of **PDGF** -C messenger RNA in the metanephric mesenchyme during epithelial conversion. Analysis of kidneys lacking the **PDGF** alpha-receptor shows selective loss of mesenchymal cells adjacent to sites of expression of **PDGF** -C mRNA; this is not found in kidneys from animals lacking **PDGF** -A or both **PDGF** -A and **PDGF** -B, indicating that **PDGF** -C may have a unique function.

MEDICAL DESCRIPTORS:

receptor binding; enzyme activity; cell proliferation; fibroblast;  
**transgenic mouse** ; in situ hybridization; nonhuman; mouse; animal experiment; controlled study; animal cell; article; priority journal

6/3,K/17 (Item 6 from file: 73)

DIALOG(R) File 73:EMBASE

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10815692 EMBASE No: 2000295686

**Platelet-derived growth factor-A-induced retinal gliosis protects against ischemic retinopathy**

Yamada H.; Yamada E.; Ando A.; Seo M.-S.; Esumi N.; Okamoto N.; Vinores M.; LaRochelle W.; Zack D.J.; Campochiaro P.A.

Dr. P.A. Campochiaro, Maumenee 719, Johns Hopkins Univ. Sch. of Medicine, 600 N. Wolfe Street, Baltimore, MD 21287-9277 United States

AUTHOR EMAIL: pcampo@jhmi.edu

American Journal of Pathology ( AM. J. PATHOL. ) (United States) 2000, 156/2 (477-487)

CODEN: AJPAA ISSN: 0002-9440

DOCUMENT TYPE: Journal; Article

LANGUAGE: ENGLISH SUMMARY LANGUAGE: ENGLISH

NUMBER OF REFERENCES: 38

...along retinal blood vessels and have been hypothesized to participate in the induction and maintenance of the blood-retinal barrier. Platelet-derived growth factor-A ( **PDGF** -A) is normally produced by retinal ganglion cells and is involved in astrocyte recruitment and proliferation. We used gain-of-function **transgenic** mice that express **PDGF** -A in photoreceptors to explore the roles of **PDGF** -A and astrocytes in the retina. Transgene-positive mice developed glial infiltration of the inner retina and had significantly less oxygen-induced retinal vascular closure and no neovascularization compared with littermate controls, which had prominent vascular closure and neovascularization. The increased survival of endothelial cells in **transgenic** mice in the face of oxygen-induced down-regulation of vascular endothelial growth factor was accompanied by an increase in astrocyte-derived fibroblast growth factor-2. Therefore, **PDGF** -A increases retinal astrocytes, which promote the survival of endothelial

cells as well as their expression of barrier characteristics.

MEDICAL DESCRIPTORS:

**transgenic mouse** ; photoreceptor; astrocyte; cell survival; endothelium cell; blood retina barrier; nonhuman; mouse; animal model; controlled study ; animal tissue; article; priority journal

6/3,K/18 (Item 7 from file: 73)

DIALOG(R)File 73:EMBASE

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07723280 EMBASE No: 1999199651

**Emphysematous lesions, inflammation, and fibrosis in the lungs of transgenic mice overexpressing platelet-derived growth factor**

Hoyle G.W.; Li J.; Finkelstein J.B.; Eisenberg T.; Liu J.-Y.; Lasky J.A.; Athas G.; Morris G.F.; Brody A.R.

Dr. G.W. Hoyle, Section of Pulmonary Diseases, Critical Care and Env'tl. Medicine, Tulane University Medical Center, 1430 Tulane Avenue, New Orleans, LA 70112 United States

AUTHOR EMAIL: ghoyle@tmc.tulane.edu

American Journal of Pathology ( AM. J. PATHOL. ) (United States) 1999, 154/6 (1763-1775)

CODEN: AJPAA ISSN: 0002-9440

DOCUMENT TYPE: Journal; Article

LANGUAGE: ENGLISH SUMMARY LANGUAGE: ENGLISH

NUMBER OF REFERENCES: 58

Because of its expression pattern and its potent effects on mesenchymal cells, platelet-derived growth factor ( **PDGF** ) has been implicated as an important factor in epithelial-mesenchymal cell interactions during normal lung development and in the pathogenesis of fibrotic lung disease. To further explore the role of **PDGF** in these processes, we have developed

**transgenic** mice that express the **PDGF** -B gene from the lung-specific surfactant protein C (SPC) promoter. Adult SPC-PDGFB **transgenic** mice exhibited lung pathology characterized by enlarged airspaces, inflammation, and fibrosis. Emphysematous changes frequently occurred throughout the lung, but inflammation and fibrotic lesions were usually confined to focal areas. The severity of this phenotype varied significantly among individual mice within the same SPC-PDGFB **transgenic** lineage. A pathology similar to that observed in adult mice was noted in lungs from **transgenic** mice as young as 1 week of age. Neonatal **transgenic** mice exhibited enlarged sacculles and thickened primary septa. Results of these studies indicated that overexpression of **PDGF** -B induced distinct abnormalities in the developing and adult lung and led to a complex phenotype that encompassed aspects of both emphysema and fibrotic lung...

MEDICAL DESCRIPTORS:

inflammatory disease; **transgenic mouse** ; protein expression; gene overexpression; phenotype; disease severity; histopathology; nonhuman; mouse; animal model; controlled study; animal tissue; article; priority journal

6/3,K/19 (Item 8 from file: 73)

DIALOG(R)File 73:EMBASE

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07639490 EMBASE No: 1999117351

**A vascular bed-specific pathway regulates cardiac expression of endothelial nitric oxide synthase**

Guillot P.V.; Guan J.; Liu L.; Kuivenhoven J.A.; Rosenberg R.D.; Sessa

W.C.; Aird W.C.

W.C. Aird, Division of Molecular Medicine, Beth Israel Deaconess Medical School, Boston, MA 02215 United States

AUTHOR EMAIL: waird@bidmc.harvard.edu

Journal of Clinical Investigation ( J. CLIN. INVEST. ) (United States)

15 MAR 1999, 103/6 (799-805)

CODEN: JCINA ISSN: 0021-9738

DOCUMENT TYPE: Journal; Article

LANGUAGE: ENGLISH SUMMARY LANGUAGE: ENGLISH

NUMBER OF REFERENCES: 50

...a variety of extracellular signals under both in vitro and in vivo conditions. To gain insight into the mechanisms underlying environmental regulation of eNos expression, **transgenic** mice were generated with the 1,600-bp 5' flanking region of the human enos promoter coupled to the coding region of the LacZ gene...

...by conditioned media from cardiac myocytes, skeletal myocytes, and brain astrocytes. Cardiac myocyte-mediated induction was partly abrogated by neutralizing anti-platelet-derived growth factor ( **PDGF** ) antibodies. In addition, promoter activity was upregulated by **PDGF** -AB. Analysis of promoter deletions revealed that a **PDGF** response element lies between -744 and -1,600 relative to the start site of transcription, whereas a **PDGF** -independent cardiac myocyte response element is present within the first 166 bp of the 5' flanking region. Taken together, these results suggest that the eNos gene is regulated in the cardiac endothelium by both a **PDGF** -dependent and **PDGF** - independent microvascular bed-specific signaling pathway.

MEDICAL DESCRIPTORS:

protein expression; enzyme regulation; **transgenic mouse** ; promoter region; culture medium; enzyme induction; DNA flanking region; signal transduction; nonhuman; mouse; animal cell; article; priority journal

6/3,K/20 (Item 9 from file: 73)

DIALOG(R)File 73:EMBASE

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07239835 EMBASE No: 1998138327

**The APP and PS1/2 mutations linked to early onset familial Alzheimer's disease increase the extracellular concentration of A/beta1-42 (43)**

Younkin S.G.

Dr. S.G. Younkin, Department of Pharmacology, Mayo Clinic, Jacksonville, FL United States

Clinical Neurology ( CLIN. NEUROL. ) (Japan) 1997, 37/12 (1099)

CODEN: RISHD ISSN: 0009-918X

DOCUMENT TYPE: Journal; Conference Paper

LANGUAGE: JAPANESE SUMMARY LANGUAGE: ENGLISH

...the AD state. To determine whether presenilin mutations act as true dominants, we collaborated with others to analyze Abeta1-40 and Abeta1-42 (43) in **transgenic** mice and transfected cells expressing wild type and mutant human presenilin transgenes under the control of the platelet-derived growth factor ( **PDGF** ) promoter. This analysis showed that expression of mutant, but not wild type human PS1 selectively increases Abeta1-42 (43) even when the endogenous mouse PS1...

...is to increase the extracellular concentration of Abeta42 (43). The plasma data establish that these mutations increase extracellular Abeta42 (43) in vivo. The results from **transgenic** mice establish that the PS1 mutations increase Abeta1- 42 (43) in the brain. This increase in Abeta1-42

(43) caused by the FAD-linked mutations...

MEDICAL DESCRIPTORS:

onset age; chromosome 14; chromosome 1; fibroblast; **transgenic mouse** ;  
protein expression; pathogenesis; human; nonhuman; mouse; animal experiment  
; animal model; controlled study; human cell; animal cell; conference paper

6/3,K/21 (Item 10 from file: 73)

DIALOG(R)File 73:EMBASE

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07199007 EMBASE No: 1998095986

**Transgenic hypertensive rats how a reduced angiotensin II induced (Casup 2sup +)(i) response in glomerular mesangial cells**

Tepel M.; Heidenreich S.; Zidek W.

M. Tepel, Universitätsklinik Marienhospital, Medizinische Klinik 1,  
Ruhr-Universität-Bochum, Holkeskampring 40, D-44625 Herne Germany

Life Sciences ( LIFE SCI. ) (United States) 28 NOV 1997, 62/1 (69-76)

CODEN: LIFSA ISSN: 0024-3205

PUBLISHER ITEM IDENTIFIER: S0024320597010394

DOCUMENT TYPE: Journal; Article

LANGUAGE: ENGLISH SUMMARY LANGUAGE: ENGLISH

NUMBER OF REFERENCES: 29

The effect of angiotensin II (Ang II) induced changes of cytosolic free calcium concentration ((Casup 2sup +)(i)) and growth response were investigated in **transgenic** TGR(mREN2)27 rats, a strain showing fulminant hypertension after the mouse Ren-2d renin gene has been integrated into its genome, in age- matched...

...arginine vasopressin or endothelin induced (Casup 2sup +)(i) increase were not significantly different in MC from TGR(mREN2)27 and SD. The Ang II or **PDGF** induced sup 3H-thymidine incorporation was not significantly different in MC from TGR(mREN2)27 and SD, indicating that the early growth response to Ang...

MEDICAL DESCRIPTORS:

calcium cell level; mesangium cell; glomerulus; **transgenic animal** ;  
renin angiotensin aldosterone system; spontaneously hypertensive rat;  
nonhuman; male; rat; animal experiment; animal model; controlled study;  
article

6/3,K/22 (Item 11 from file: 73)

DIALOG(R)File 73:EMBASE

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06745044 EMBASE No: 1997026520

**Lens-specific expression of PDGF-A alters lens growth and development**

Reneker L.W.; Overbeek P.A.

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Baylor Plaza, Houston, TX 77030 United States

AUTHOR EMAIL: lreneker@condor.bcm.tmc.edu

Developmental Biology ( DEV. BIOL. ) (United States) 1996, 180/2  
(554-565)

CODEN: DEBIA ISSN: 0012-1606

DOCUMENT TYPE: Journal; Article

LANGUAGE: ENGLISH SUMMARY LANGUAGE: ENGLISH

NUMBER OF REFERENCES: 29

...study the molecular mechanisms by which growth factors influence

development decisions. In this study, we have investigated the expression patterns of platelet-derived growth factor ( **PDGF** ) and **PDGF** receptors during murine eye development by in situ hybridization. Postnatally, **PDGF** -A is highly expressed in the iris and ciliary body, the ocular tissues closest to the germinative zone of the lens, a region where most proliferation of lens epithelial cells occurs. **PDGF** -A is also present in the corneal endothelium anterior to the lens epithelium in embryonic and early postnatal eyes. **PDGF** -B is expressed in the iris and ciliary body as well as in the vascular cells which surround the lens during early eye development. In the lens, expression of **PDGF** -alpha receptor ( **PDGF** -alphaR), a receptor that can bind both **PDGF** -A and **PDGF** -B, is restricted to the lens epithelium throughout life. The expression of **PDGF** -alphaR in the lens epithelial cells and **PDGF** (A- and B-chains) in the ocular tissues adjacent to the lens suggests that **PDGF** signaling may play a key role in regulating lens development. To further examine how **PDGF** affects lens development in vivo, we generated **transgenic** mice that express human **PDGF** -A in the lens under the control of the alphaA-crystallin promoter. The **transgenic** mice exhibit lenticular defects that result in cataracts. The percentage of surface epithelial cells in S-phase is increased in **transgenic** lenses compared to their nontransgenic littermates. Higher than normal levels of cyclin A and cyclin D2 expression were also detected in **transgenic** lens epithelium. These results together suggest that **PDGF** -A can induce a proliferative response in lens epithelial cells. The lens epithelial cells in the **transgenic** mice also exhibit characteristics of differentiating fiber cells. For example, the **transgenic** lens epithelial cells are slightly elongated, contain larger and less condensed nuclei, and express fiber-cell-specific beta-crystallins. Our results suggest that **PDGF** -A normally acts as a proliferative factor for the lens epithelial cells in vivo. Elevated levels of **PDGF** -A enhance proliferation, but also appear to induce some aspects of the fiber cell differentiation pathway.

MEDICAL DESCRIPTORS:

animal experiment; animal tissue; article; embryo; immunohistochemistry; in situ hybridization; mouse; nonhuman; priority journal; **transgenic mouse**

6/3,K/23 (Item 12 from file: 73)

DIALOG(R)File 73:EMBASE

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06734245 EMBASE No: 1997015715

**PDGF mediates a neuron-astrocyte interaction in the developing retina**

Fruttiger M.; Calver A.R.; Kruger W.H.; Mudhar H.S.; Michalovich D.; Takakura N.; Shin Ichi Nishikawa; Richardson W.D.

M. Fruttiger, MRC Lab. for Molecular Cell Biology, Department of Biology, University College London, London WC1E 6BT United Kingdom

Neuron ( NEURON ) (United States) 1996, 17/6 (1117-1131)

CODEN: NERNE ISSN: 0896-6273

DOCUMENT TYPE: Journal; Article

LANGUAGE: ENGLISH SUMMARY LANGUAGE: ENGLISH

NUMBER OF REFERENCES: 75

...the developing retina from the optic nerve head, over the axons of retinal ganglion cells (RGCs). RGCs express the platelet- derived growth factor A-chain ( **PDGF** -A) and retinal astrocytes the **PDGF** alpha- receptor (PDGFRalpha), suggesting that **PDGF** mediates a paracrine interaction between these cells. To test this, we inhibited **PDGF** signaling in the eye with a neutralizing anti-PDGFRalpha antibody or a soluble extracellular fragment of PDGFRalpha. These treatments inhibited development of the astrocyte network. We also generated **transgenic** mice that overexpress

**PDGF** -A in RGCs. This resulted in hyperproliferation of astrocytes, which in turn induced excessive vasculogenesis. Thus, **PDGF** appears to be a link in the chain of cell-cell interactions responsible for matching numbers of neurons, astrocytes, and blood vessels during retinal development.

MEDICAL DESCRIPTORS:

...animal cell; animal model; animal tissue; article; astrocyte; cell proliferation; controlled study; monkey; nerve cell; nonhuman; optic nerve; priority journal; retina ganglion cell; retina neovascularization; **transgenic mouse**

6/3,K/24 (Item 13 from file: 73)

DIALOG(R)File 73:EMBASE

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06664785 EMBASE No: 1996329666

**Fibroblast growth factor receptor 1-induced differentiation of endothelial cell line established from tsA58 large T transgenic mice**

Kanda S.; Landgren E.; Ljungstrom M.; Claesson-Welsh L.

Biomedical Center, Ludwig Institute for Cancer Research, Box 595,S-751 24 Uppsala Sweden

Cell Growth and Differentiation ( CELL GROWTH DIFFER. ) (United States) 1996, 7/3 (383-395)

CODEN: CGDIE ISSN: 1044-9523

DOCUMENT TYPE: Journal; Article

LANGUAGE: ENGLISH SUMMARY LANGUAGE: ENGLISH

...basement membrane, migration, proliferation, and differentiation. To study differentiation of endothelial cells, we established a brain capillary endothelial cell line from H-2Ksup b- tsA58 **transgenic** mice. These cells are stable at 33degreeC and display endothelial cell-specific characters, such as expression of von Willebrand factor and binding sites for the...

...panel of growth factors on cellular responses. A number of factors, such as hepatocyte growth factor, vascular endothelial growth factor, and platelet-derived growth factor ( **PDGF** )-AA failed to induce biological responses. **PDGF** -BB, epidermal growth factor, and acidic and basic fibroblast growth factor (FGF) induced proliferation of the cells. Of all the factors tested, only acidic FGF...

...their effects on plasminogen activator (PA)-induction and migration of the cells. Transfected cells, expressing a chimeric receptor, composed of the extracellular part from the **PDGF** alpha-receptor and the intracellular part from FGF receptor-1, responded to **PDGF** -AA treatment with plasminogen activator induction, migration, proliferation, and tube formation in collagen. These results indicate that FGF receptor-1 coupled to signal transduction pathways...

MEDICAL DESCRIPTORS:

animal cell; article; controlled study; dna transfection; mouse; nonhuman; priority journal; protein expression; **transgenic mouse** ; vascular endothelium

6/3,K/25 (Item 14 from file: 73)

DIALOG(R)File 73:EMBASE

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06317400 EMBASE No: 1995354830

**Levels and alternative splicing of amyloid beta protein precursor (APP)**

**transcripts in brains of APP transgenic mice and humans with Alzheimer's disease**

Rockenstein E.M.; McConlogue L.; Tan H.; Power M.; Masliah E.; Mucke L.  
 Gladstone/Neurology Neurobiol. Prog., Gladstone Institutes, P. O. Box  
 419100, San Francisco, CA 94141-9100 United States  
 Journal of Biological Chemistry ( J. BIOL. CHEM. ) (United States) 1995  
 , 270/47 (28257-28267)  
 CODEN: JBCHA ISSN: 0021-9258  
 DOCUMENT TYPE: Journal; Article  
 LANGUAGE: ENGLISH SUMMARY LANGUAGE: ENGLISH

Abnormal expression of human amyloid precursor protein (hAPP) gene products may play a critical role in Alzheimer's disease (AD). Recently, a **transgenic** model was established in which platelet-derived growth factor ( **PDGF** ) promoter-driven neuronal expression of an alternatively spliced hAPP minigene resulted in prominent AD-type neuropathology (Games, D., Adams, D., Alessandrini, R., Barbour, R., Berthelette...

...L., and Penniman, E. (1995) Nature 373, 523-527). Here we compared the levels and alternative splicing of APP transcripts in brain tissue of hAPP **transgenic** and nontransgenic mice and of humans with and without AD. **PDGF** -hAPP mice showed severalfold higher levels of total APP mRNA than did nontransgenic mice or humans, whereas their endogenous mouse APP mRNA levels were decreased...

...resulted in a high ratio of mRNAs encoding mutated hAPP versus wild-type mouse APP. Modifications of hAPP introns 6, 7, and 8 in the **PDGF** -hAPP construct resulted in a prominent change in alternative splice site selection with transcripts encoding hAPP770 or hAPP751 being expressed at substantially higher levels than...

**MEDICAL DESCRIPTORS:**

alternative rna splicing; animal tissue; article; frontal cortex; gene expression; human; human tissue; intron; nonhuman; priority journal; rna analysis; **transgenic mouse**

**6/3,K/26 (Item 15 from file: 73)**

DIALOG(R) File 73:EMBASE

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06118472 EMBASE No: 1995149206

**Luteal failure in transgenic mice carrying a PDGF□dominant-negative□ mutant/GH hybrid transgene**

Pekny M.; Pekna M.; Ostman A.; Tornell J.; Feinstein R.; Forsberg-Nilsson K.; Heldin C.-H.; Westermarck B.; Betsholtz C.  
 Department of Medical Biochemistry, University of Goteborg,  
 Medicinaregatan 9,S-413 90 Goteborg Sweden  
 Transgenics ( TRANSGENICS ) (United Kingdom) 1995, 1/5 (515-523)  
 CODEN: TADTE ISSN: 1023-6171  
 DOCUMENT TYPE: Journal; Article  
 LANGUAGE: ENGLISH SUMMARY LANGUAGE: ENGLISH

**Luteal failure in transgenic mice carrying a PDGF□dominant-negative□ mutant/GH hybrid transgene**

**MEDICAL DESCRIPTORS:**

\*luteal insufficiency; \* **transgenic mouse**

**6/3,K/27 (Item 16 from file: 73)**

DIALOG(R) File 73:EMBASE

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04569154 EMBASE No: 1991063197

**PDGF B-chain in neurons of the central nervous system, posterior pituitary, and in a transgenic model**

Sasahara M.; Fries J.W.U.; Raines E.W.; Gown A.M.; Westrum L.E.; Frosch M.P.; Bonthron D.T.; Ross R.; Collins T.

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United States

Cell ( CELL ) (United States) 1991, 64/1 (217-227)

CODEN: CELLB ISSN: 0092-8674

DOCUMENT TYPE: Journal; Article

LANGUAGE: ENGLISH SUMMARY LANGUAGE: ENGLISH

**PDGF B-chain in neurons of the central nervous system, posterior pituitary, and in a transgenic model**

...regulatory molecules that stimulate chemotaxis, proliferation, and increased metabolism of primarily connective tissue cells. In a survey of normal tissues, we found specific immunostaining for **PDGF** B-chain in neurons, principal dendrites, some axons, and probable terminals throughout the brain, in the dorsal horn of the spinal cord, and in the posterior pituitary of a nonhuman primate (*Macaca nemestrina*). **PDGF** activity was extracted from brain cortex and posterior pituitary, and ubiquitous expression of transcripts for the two chains of **PDGF** and both **PDGF** receptors was detected throughout the brain and posterior pituitary. A **transgenic** model was also evaluated in which the chloramphenicol acetyltransferase gene was placed under transcriptional control of the **PDGF** B-chain promoter. The transgene was preferentially expressed within neural cell bodies in the cortex, hippocampus, and cerebellum. **PDGF** may act as a neuronal regulatory agent. Neuronal release of **PDGF** could contribute to nerve regeneration and to glial proliferation that leads to gliosis and scarring.

**MEDICAL DESCRIPTORS:**

\*central nervous system; \*gene expression; \*neurohypophysis; \* **transgenic animal**

?

Set	Items	Description
S1	292	(PDGF OR PDGF-AA OR PDGF-AB OR PDGF-BB) (S) (MILK OR TRANS-GENIC OR BIOREACTOR)
S2	2123	(PROMOTER) (S) (LACTALBUMIN OR CASEIN OR LACTOGLOBULIN OR - (MAMMARY (W) EPITHELIAL))
S3	0	S1 AND S2
S4	82	S1 AND (TRANSGENIC (W) (ANIMAL OR MOUSE OR RAT OR BOVINE OR GOAT OR PIG))
S5	31	S4 NOT PY>2000
S6	27	RD (unique items)

?

**COST**

05apr06 09:48:58 User259876 Session D860.2  
\$1.78 0.523 DialUnits File155  
\$0.66 3 Type(s) in Format 3  
\$0.66 3 Types  
\$2.44 Estimated cost File155  
\$4.00 0.678 DialUnits File5  
\$1.28 8 Type(s) in Format 95 (KWIC)  
\$1.28 8 Types

\$5.28 Estimated cost File5  
\$7.47 0.667 DialUnits File73  
\$49.60 16 Type(s) in Format 3  
\$49.60 16 Types  
\$57.07 Estimated cost File73  
OneSearch, 3 files, 1.868 DialUnits FileOS  
\$1.33 INTERNET  
\$66.12 Estimated cost this search  
\$66.98 Estimated total session cost 2.098 DialUnits

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